FORM PTO-1390 (Modified) (REV 11-98) U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371

ATTORNEY'S DOCKET NUMBER	U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR
8830-8	09/937687

INTERNATIONAL APPLICATION NO.	INTERNATIONAL FILING DATE	PRIORITY DATE CLAIMED
PCT/GB00/01089	March 29, 2000	March 29, 1999

TITLE OF INVENTION
Peptide

APPLICANT(S) FOR DO/EO/US
Finbarr Paul Mary O'Harte and Peter Raymond Flatt

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c) (2))
   a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau).
   b. ☐ has been transmitted by the International Bureau.
   c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ A copy of the International Search Report (PCT/ISA/210).
8. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
   a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
   b. ☐ have been transmitted by the International Bureau.
   c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
   d. ☒ have not been made and will not be made.
9. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
10. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
11. ☐ A copy of the International Preliminary Examination Report (PCT/IPEA/409).
12. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).

Items 13 to 20 below concern document(s) or information included:

13. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
14. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
15. ☒ A **FIRST** preliminary amendment.
16. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
17. ☐ A substitute specification.
18. ☐ A change of power of attorney and/or address letter.
19. ☒ Certificate of Mailing by Express Mail
20. ☒ Other items or information:

Unexecuted Declaration and Power of Attorney
Express Mail Label No. EL 813776259 US
Small Entity Status Is Claimed

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR)	INTERNATIONAL APPLICATION NO.	ATTORNEY'S DOCKET NUMBER
09/937687	PCT/GB00/01089	8830-8

21. The following fees are submitted:.

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :		CALCULATIONS PTO USE ONLY
☐ Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2) paid to USPTO and International Search Report not prepared by the EPO or JPO	$1,000.00	
☒ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but Internation Search Report prepared by the EPO or JPO	$860.00	
☐ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO	$710.00	
☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4)	$690.00	
☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4)	$100.00	
ENTER APPROPRIATE BASIC FEE AMOUNT =		$860.00
Surcharge of **$130.00** for furnishing the oath or declaration later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).		$0.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total claims	17 - 20 =	0	x $18.00	$0.00	
Independent claims	3 - 3 =	0	x $80.00	$0.00	
Multiple Dependent Claims **(check if applicable)**.			☒	$270.00	
TOTAL OF ABOVE CALCULATIONS			=	$1,130.00	
Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28) **(check if applicable)**.			☒	$565.00	
SUBTOTAL			=	$565.00	
Processing fee of **$130.00** for furnishing the English translation later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).			+	$0.00	
TOTAL NATIONAL FEE			=	$565.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) **(check if applicable).**			☐	$0.00	
TOTAL FEES ENCLOSED			=	$565.00	
				Amount to be: refunded	$
				charged	$

☒ A check in the amount of **$565.00** to cover the above fees is enclosed.

☐ Please charge my Deposit Account No. in the amount of to cover the above fees.
A duplicate copy of this sheet is enclosed.

☒ The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account No. **50-0573** A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

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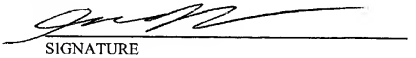

SIGNATURE

DANIEL A. MONACO
NAME

30,480
REGISTRATION NUMBER

September 28, 2001
DATE

09/937687

CERTIFICATE OF MAILING BY "EXPRESS MAIL" (37 CFR 1.10) Applicant(s): **O'Harte et al.**			Docket No. **8830-8**
Serial No. **PCT/GB00/01089**	Filing Date **Int'l 3/29/00**	Examiner	Group Art Unit

Invention:

Peptide

I hereby certify that this **US Entry into National Phase of PCT/GB00/01089 and formal documents**
(Identify type of correspondence)

is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 in an envelope addressed to: The Commissioner of Patents and Trademarks, Washington, D.C. 20231-0001 on **September 28, 2001**
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PATENT

Attorney Docket No.: 8830-8

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re:	Patent application of Finbarr Paul Mary O'Harte et al.	:	
Serial No.:	(International Application PCT/GB00/01089)	:	Group Art Unit:
Filed:	(International Application: March 29, 2000)	:	Examiner:
For:	Peptide	:	

Preliminary Amendment

Commissioner for Patents
Washington, D.C. 20231

Sir:

Kindly amend the above-identified patent application, prior to calculation of the filing fee, as follows.

In the Specification

Insert the Abstract attached hereto as a separate page.

In the Claims

CERTIFICATE OF MAILING
UNDER 37 C.F.R. 1.10

EXPRESS MAIL Mailing Label Number: EL 813776259 US
Date of Deposit: **September 28, 2001**
I hereby certify that this correspondence, along with any paper referred to as being attached or enclosed, and/or fee, is being deposited with the United States Postal Service, "EXPRESS MAIL-POST OFFICE TO ADDRESSEE" service under 37 CFR 1.10, on the date indicated above, and addressed to: Commissioner for Patents, Washington, D.C. 20231.

Signature of person mailing page:

Therese McKinley
Type or print name of person

Add the following new claim 12:

12. (new) A method for treating diabetes comprising administering to an individual in need of such treatment an effective amount of an analog according to claim 1 or 3.

Rewrite claims 4-6 and 8 to read as follows. A mark-up of the amended claims is submitted herein as Appendix A.

4. (amended) A peptide analogue as claimed in claim 1 or 3 wherein the substitution or modification is chosen from the group comprising D-amino acid substitutions in 1, 2 and/or 3 positions and/or N terminal glycation, alkylation, acetylation or acylation.

5. (amended) A peptide analogue as claimed in claim 1 or 3 wherein the amino acid in the 2 or 3 position is substituted by lysine, serine, 4-amino butyric, Aib, D-alanine, Sarcosine or Proline.

6. (amended) An analogue as claimed in claim 1 or 3 wherein the N terminus is modified by one of the group of modifications including glycation, alkylation, acetylation or by the addition of an isopropyl group.

8. (amended) A pharmaceutical composition including an analogue as claimed in claim 1 or 3.

Remarks

Claims 1-6 and 8-12 are pending in the application. The dependencies of certain claims have been reduced, to conform to United States practice. Claim 7 has been cancelled and presented as claim 12, in a method of treatment format consistent with United States practice. No new matter has been introduced.

An Abstract is submitted herewith, which is identical to the abstract appearing in the international application.

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APEENDIX A: Mark-up of amended claims

4. (amended) A peptide analogue as claimed in [any of the preceding claims] claim 1 or 3 wherein the substitution or modification is chosen from the group comprising D-amino acid substitutions in 1, 2 and/or 3 positions and/or N terminal glycation, alkylation, acetylation or acylation.

5. (amended) A peptide analogue as claimed in [any of the preceding claims] claim 1 or 3 wherein the amino acid in the 2 or 3 position is substituted by lysine, serine, 4-amino butyric, Aib, D-alanine, Sarcosine or Proline.

6. (amended) An analogue as claimed in [any of the preceding claims] claim 1 or 3 wherein the N terminus is modified by one of the group of modifications [include] including glycation, alkylation, acetylation or by the addition of an isopropyl group.

8. (amended) A pharmaceutical composition including an analogue as claimed in [any of the preceding claims] claim 1 or 3.

Abstract

The present invention provides peptides which stimulate the release of insulin. The peptides, based on GIP 1-42, include substitutions and/or modifications which enhance and influence secretion and/or have enhanced resistance to degradation. The invention also provides a process of N terminally modifying GIP and the use of the peptide analogues for treatment of diabetes.

"Peptide"

The present invention relates to the release of insulin and the control of blood glucose concentration. More particularly the invention relates to the use of peptides to stimulate release of insulin, lowering of blood glucose and pharmaceutical preparations for treatment of type 2 diabetes.

Gastric inhibitory polypeptide (GIP) and glucagon-like peptide-1(7-36)amide (truncated GLP-1; tGLP-1) are two important insulin-releasing hormones secreted from endocrine cells in the intestinal tract in response to feeding. Together with autonomic nerves they play a vital supporting role to the pancreatic islets in the control of blood glucose homeostasis and nutrient metabolism.

Dipeptidyl peptidase IV (DPP IV; EC 3.4.14.5) has been identified as a key enzyme responsible for inactivation of GIP and tGLP-1 in serum. DPP IV is completely inhibited in serum by the addition of diprotin A(DPA, 0.1 mmol/l). This occurs through the rapid removal of

the N-terminal dipeptides Tyr^1- Ala^2 and His^7-Ala^8 giving rise to the main metabolites GIP(3-42) and GLP-1(9-36)amide, respectively. These truncated peptides are reported to lack biological activity or to even serve as antagonists at GIP or tGLP-1 receptors. The resulting biological half-lives of these incretin hormones *in vivo* are therefore very short, estimated to be no longer than 5 min.

In situations of normal glucose regulation and pancreatic B-cell sensitivity, this short duration of action is advantageous in facilitating momentary adjustments to homeostatic control. However, the current goal of a possible therapeutic role of incretin hormones, particularly tGLP-1 in NIDDM therapy is frustrated by a number of factors in addition to finding a convenient route of administration. Most notable of these are rapid peptide degradation and rapid absorption (peak concentrations reached 20 min) and the resulting need for both high dosage and precise timing with meals. Recent therapeutic strategies have focused on precipitated preparations to delay peptide absorption and inhibition of GLP-1 degradation using specific inhibitors of DPP IV. A possible therapeutic role is also suggested by the observation that a specific inhibitor of DPP IV, isoleucine thiazolidide, lowered blood glucose and enhanced insulin secretion in glucose-treated diabetic obese Zucker rats presumably by protecting against catabolism of the incretin hormones tGLP-1 and GIP.

Numerous studies have indicated that tGLP-1 infusion restores pancreatic B-cell sensitivity, insulin secretory oscillations and improved glycemic control in various groups of patients with IGT or NIDDM. Longer term studies also show significant benefits of tGLP-1 injections in NIDDM and possibly IDDM therapy, providing a major incentive to develop an orally effective or long-acting tGLP-1 analogue. Several attempts have been made to produce structurally modified analogues of tGLP-1 which are resistant to DPP IV degradation. A significant extension of serum half-life is observed with His^7- glucitol tGLP-1 and tGLP-1 analogues substituted at position 8 with Gly, Aib, Ser or Thr. However, these structural modifications seem to impair receptor binding and insulinotrophic activity thereby compromising part of the benefits of protection from proteolytic degradation. In recent studies using His^7-glucitol tGLP-1, resistance to DPP IV and serum degradation was accompanied by severe loss of insulin-releasing activity.

GIP shares not only the same degradation pathway as tGLP-1 but many similar physiological actions, including stimulation of insulin and somatostatin secretion, and the enhancement of glucose disposal. These actions are viewed as key aspects in the antihyperglycemic properties of tGLP-1, and there is therefore good expectation that GIP may have similar potential as NIDDM therapy. Indeed, compensation by GIP is held to explain the modest disturbances of glucose homeostasis observed in tGLP-1 knockout mice. Apart from early studies, the anti-diabetic potential

of GIP has not been explored and tGLP-1 may seem more attractive since it is viewed by some as a more potent insulin secretagogue when infused at "so called" physiological concentrations estimated by RIA.

The present invention aims to provide effective analogues of GIP. It is one aim of the invention to provide a pharmaceutical for treatment of Type 2 diabetes.

According to the present invention there is provided an effective peptide analogue of the biologically active GIP(1-42) which has improved characteristics for treatment of Type 2 diabetes wherein the analogue comprises at least 15 amino acid residues from the N terminus of GIP(1-42) and has at least one amino acid substitution or modification at position 1-3 and not including Tyr^1 glucitol GIP(1-42).

The structures of human and porcine GIP(1-42) are shown below. The porcine peptide differs by just two amino acid substitutions at positions 18 and 34.

The analogue may include modification by fatty acid addition at an epsilon amino group of at least one lysine residue.

The invention includes Tyr^1 glucitol GIP(1-42) having fatty acid addition at an epsilon amino group of at least one lysine residue.

Fig. 1. Primary structure of human gastric inhibitory polypeptide (GIP)

1 5 10 15 20 25
NH_2-Tyr-Ala-Glu-Gly-Thr-Phe-Ile-Ser-Asp-Tyr-Ser-Ile-Ala-Met-Asp-Lys-Ile-His-Gln-Gln-Asp-Phe-Val-Asn-Trp-

Leu-Leu-Ala-Gln-Lys-Gly-Lys-Lys-Asn-Asp-Trp-Lys-His-Asn-Ile-Thr-Gln-COOH
30 35 40

Fig. 2. Primary structure of porcine gastric inhibitory polypeptide (GIP)

1 5 10 15 20 25
NH_2-Tyr-Ala-Glu-Gly-Thr-Phe-Ile-Ser-Asp-Tyr-Ser-Ile-Ala-Met-Asp-Lys-Ile-Arg-Gln-Gln-Asp-Phe-Val-Asn-Trp-

Leu-Leu-Ala-Gln-Lys-Gly-Lys-Lys-Ser-Asp-Trp-Lys-His-Asn-Ile-Thr-Gln-COOH
30 35 40

Analogues of GIP(1-42) may have an enhanced capacity to stimulate insulin secretion, enhance glucose disposal, delay glucose absorption or may exhibit enhanced stability in plasma as compared to native GIP. They also may have enhanced resistance to degradation.

Any of these properties will enhance the potency of the analogue as a therapeutic agent.

Analogues having D-amino acid substitutions in the 1, 2 and 3 positions and/or N-glycated, N-alkylated, N-acetylated or N-acylated amino acids in the 1 position are resistant to degradation *in vivo*.

Various amino acid substitutions at second and third amino terminal residues are included, such as GIP(1-42)Gly2, GIP(1-42)Ser2, GIP(1-42)Abu2, GIP(1-42)Aib, GIP(1-42)D-Ala2, GIP(1-42)Sar2, and GIP(1-42)Pro3.

Amino-terminally modified GIP analogues include N-glycated GIP(1-42), N-alkylated GIP(1-42), N-actylated

GIP(1-42), N-acetyl-GIP(1-42) and N-isopropyl GIP(1-42).

Other stabilised analogues include those with a peptide isostere bond between amino terminal residues at position 2 and 3. These analogues may be resistant to the plasma enzyme dipeptidyl-peptidase IV (DPP IV) which is largely responsible for inactivation of GIP by removal of the amino-terminal dipeptide Tyrl-Ala2.

In particular embodiments, the invention provides a peptide which is more potent than human or porcine GIP in moderating blood glucose excursions, said peptide consisting of GIP(1-42) or N-terminal fragments of GIP(1-42) consisting of up to between 15 to 30 amino acid residues from the N-terminus (i.e. GIP(1-15) - GIP(1-3)) with one or more modifications selected from the group consisting of:

(a) substitution of Ala^2 by Gly
(b) substitution of Ala^2 by Ser
(c) substitution of Ala^2 by Abu
(d) substitution of Ala^2 by Aib
(e) substitution of Ala^2 by D-Ala
(f) substitution of Ala^2 by Sar
(g) substitution of Glu^3 by Pro
(h) modification of Tyr^1 by acetylation
(i) modification of Tyr^1 by acylation
(j) modification of Tyr^1 by alkylation
(k) modification of Tyr^1 by glycation
(l) conversion of Ala^2-Glu^3 bond to a psi [CH2NH] bond

(m) conversion of Ala2-Glu3 bond to a stable peptide isotere bond

(n) (n-isopropyl-H) 1GIP.

The invention also provides the use of Tyr^1-glucitol GIP in the preparation of a medicament for the treatment of diabetes.

The invention further provides improved pharmaceutical compositions including analogues of GIP with improved pharmacological properties.

Other possible analogues include certain commonly encountered amino acids, which are not encoded by the genetic code, for example, beta-alanine (beta-ala), or other omega-amino acids, such as 3-amino propionic, 4-amino butyric and so forth, ornithine (Orn), citrulline (Cit), homoarginine (Har), t-butylalanine (t-BuA), t-butylglycine (t-BuG), N-methylisoleucine (N-MeIle), phenylglycine (Phg), and cyclohexylalanine (Cha), norleucine (Nle), cysteic acid (Cya) and methionine sulfoxide (MSO), substitution of the D form of a neutral or acidic amino acid or the D form of tyrosine for tyrosine.

According to the present invention there is also provided a pharmaceutical composition useful in the treatment of diabetes type II which comprises an effective amount of the peptide as described herein, in admixture with a pharmaceutically acceptable excipient.

The invention also provides a method of N-terminally modifying GIP or analogues thereof the method comprising the steps of synthesizing the peptide from the C terminal to the penultimate N terminal amino acid, adding tyrosine to a bubbler system as a F-moc protected Tyr(tBu)-Wang resin, deprotecting the N-terminus of the tyrosine and reacting with the modifying agent, allowing the reaction to proceed to completion, cleaving the modified tyrosine from the Wang resin and adding the modified tyrosine to the peptide synthesis reaction.

Preferably the agent is glucose, acetic anhydride or pyroglutamic acid.

The invention will now be demonstrated with reference to the following non-limiting example and the accompanying figures wherein:

Figure 1a illustrates degradation of GIP by DPP IV.

Figure 1b illustrates degradation of GIP and Tyr^1 glucitol GIP by DPP IV.

Figure 2a illustrates degradation of GIP human plasma.

Figure 2b illustrates degradation of GIP and Tyr^1-glucitol GIP by human plasma.

Figure 3 illustrates electrospray ionization mass spectrometry of GIP, Tyr^1-glucitol GIP and the major degradation fragment GIP(3-42).

Figure 4 shows the effects of GIP and glycated GIP on plasma glucose homeostasis.

Figure 5 shows effects of GIP on plasma insulin responses.

Figure 6 illustrates DPP-IV degradation of GIP 1-42.

Figure 7 illustrates DPP-IV degradation of GIP (Abu^2).

Figure 8 illustrates DPP-IV degradation of GIP (Sar^2).

Figure 9 illustrates DPP-IV degradation of GIP (Ser^2),

Figure 10 illustrates DPP-IV degradation of N-Acetyl-GIP.

Figure 11 illustrates DPP-IV degradation of glycated GIP.

Figure 12 illustrates human plasma degradation of GIP.

Figure 13 illustrates human plasma degradation of GIP (Abu^2).

Figure 14 illustrates human plasma degradation of GIP (Sar^2).

Figure 15 illustrates human plasma degradation of GIP (Ser^2).

Figure 16 illustrates human plasma degradation of glycated GIP.

Figure 17 shows the effects of various concentrations of GIP 1-42 and GIP (Abu^2) on insulin release from BRIN-BD11 cells incubated at 5.6mM glucose.

Figure 18 shows the effects of various concentrations of GIP 1-42 and GIP (Abu^2) on insulin release from BRIN-BD11 cells incubated at 16.7mM glucose.

Figure 19 shows the effects of various concentrations of GIP 1-42 and GIP (Sar^2) on insulin release from BRIN-BD11 cells incubated at 5.6mM glucose.

Figure 20 shows the effects of various concentrations of GIP 1-42 and GIP (Sar^2) on insulin release from BRIN-BD11 cells incubated at 16.7mM glucose.

Figure 21 shows the effects of various concentrations of GIP 1-42 and GIP (Ser^2) on insulin release from BRIN-BD11 cells incubated at 5.6mM glucose.

Figure 22 shows the effects of various concentrations of GIP 1-42 and GIP (Ser^2) on insulin release from BRIN-BD11 cells incubated at 16.7mM glucose.

Figure 23 shows the effects of various concentrations of GIP 1-42 and N-Acetyl-GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 5.6mM glucose.

Figure 24 shows the effects of various concentrations of GIP 1-42 and N-Acetyl-GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 16.7mM glucose.

Figure 25 shows the effects of various concentrations of GIP 1-42 and glycated GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 5.6mM glucose.

Figure 26 shows the effects of various concentrations of GIP 1-42 and glycated GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 16.7mM glucose.

Figure 27 shows the effects of various concentrations of GIP 1-42 and GIP (Gly^2) on insulin release from BRIN-BD11 cells incubated at 5.6mM glucose.

Figure 28 shows the effects of various concentrations of GIP 1-42 and GIP (Gly^2) on insulin release from BRIN-BD11 cells incubated at 16.7mM glucose.

Figure 29 shows the effects of various concentrations of GIP 1-42 and GIP (Pro^3) on insulin release from BRIN-BD11 cells incubated at 5.6mM glucose.

Figure 30 shows the effects of various concentrations of GIP 1-42 and GIP (Pro^3) on insulin release from BRIN-BD11 cells incubated at 16.7mM glucose.

Example 1

Preparation of N-terminally modified GIP and analogues thereof.

The N-terminal modification of GIP is essentially a three step process. Firstly, GIP is synthesised from its C-terminal (starting from a Fmoc-Gln (Trt)-Wang resin, Novabiochem) up to the penultimate N-terminal amino-acid (Ala2) on an automated peptide synthesizer (Applied Biosystems, CA, USA). The synthesis follows standard Fmoc peptide chemistry protocols. Secondly, the N-terminal amino acid of native GIP (Tyr) is added to a manual bubbler system as a Fmoc-protected Tyr(tBu)-Wang resin. This amino acid is deprotected at its N-terminus (piperidine in DMF (20% v/v)) and allowed to react with a high concentration of glucose (glycation, under reducing conditions with sodium cyanoborohydride), acetic anhydride (acetylation), pyroglutamic acid (pyroglutamyl) etc. for up to 24 h as necessary to allow the reaction to go to completion. The completeness of reaction will be monitored using the ninhydrin test which will determine the presence of available free a-amino groups. Thirdly, (once the reaction is complete) the now structurally modified Tyr is cleaved from the wang resin (95% TFA, and 5% of the appropriate scavengers. N.B. Tyr is considered to be a problematic amino acid and may need special consideration) and the required amount of N-terminally modified-Tyr consequently added directly to the automated peptide synthesiser, which will carry on the synthesis, therby stitching the N-terminally modified-

Tyr to the a-amino of GIP(Ala2), completing the synthesis of the GIP analogue. This peptide is cleaved off the Wang resin (as above) and then worked up using the standard Buchner filtering, precipation, rotary evaporation and drying techniques.

Example 2

The following example investigates preparation of Tyr^1-glycitol GIP together with evaluation of its antihyperglycemic and insulin-releasing properties in vivo. The results clearly demonstrate that this novel GIP analogue exhibits a substantial resistance to aminopeptidase degradation and increased glucose lowering activity compared with the native GIP.

Research Design and Methods

Materials. Human GIP was purchased from the American Peptide Company (Sunnyvale, CA, USA). HPLC grade acetonitrile was obtained from Rathburn (Walkersburn, Scotland). Sequencing grade trifluoroacetic acid (TFA) was obtained from Aldrich (Poole, Dorset, UK). All other chemicals purchased including dextran T-70, activated charcoal, sodium cyanoborohydride and bovine serum albumin fraction V were from Sigma (Poole, Dorset, UK). Diprotin A (DPA) was purchased from Calbiochem-Novabiochem (UK) Ltd. (Beeston, Nottingham, UK) and rat insulin standard for RIA was obtained form Novo Industria (Copenhagen, Denmark). Reversed-phase

Sep-Pak cartridges (C-18) were purchased from Millipore-Waters (Milford, MA, USA). All water used in these experiments was purified using a Milli-Q, Water Purification System (Millipore Corporation, Milford, MA, USA).

Preparation of Tyr^1-glucitol GIP. Human GIP was incubated with glucose under reducing conditions in 10 mmol/l sodium phosphate buffer at pH 7.4 for 24 h. The reaction was stopped by addition of 0.5 mol/l acetic acid (30 µl) and the mixture applied to a Vydac (C-18)(4.6 x 250mm) analytical HPLC column (The Separations Group, Hesperia, CA, USA) and gradient elution conditions were established using aqueous/TFA and acetonitrile/TFA solvents. Fractions corresponding to the glycated peaks were pooled, taken to dryness under vacuum using an AES 1000 Speed-Vac concentrator (Life Sciences International, Runcorn, UK) and purified to homogeneity on a Supelcosil (C-8) (4.6 x 150mm) column (Supelco Inc., Poole, Dorset, UK).

Degradation of GIP and Tyr^1-glucitol GIP by DPP IV. HPLC-purified GIP or Tyr^1-glucitol GIP were incubated at 37°C with DPP-IV (5mU) for various time periods in a reaction mixture made up to 500 µl with 50 mmol/l triethanolamine-HCl, pH 7.8 (final peptide concentration 1 µmol/l). Enzymatic reactions were terminated after 0, 2, 4 and 12 hours by addition of 5 µl of 10% (v/v) TFA/water. Samples were made up to a final volume of 1.0 ml with 0.12% (v/v) TFA and stored at -20°C prior to HPLC analysis.

Degradation of GIP and Tyr^1-glucitol GIP by human plasma. Pooled human plasma (20 µl) taken from six healthy fasted human subjects was incubated at 37°C with GIP or Tyr^1-glucitol GIP (10 µg) for 0 and 4 hours in a reaction mixture made up to 500 µl, containing 50 mmol/l triethanolamine/HCL buffer pH 7.8. Incubations for 4 hours were also performed in the presence of diprotin A (5 mU). The reactions were terminated by addition of 5 µl of TFA and the final volume adjusted to 1.0 ml using 0.1% v/v TFA/water. Samples were centrifuged (13,000g, 5 min) and the supernatant applied to a C-18 Sep-Pak cartridge (Millipore-Waters) which was previously primed and washed with 0.1% (v/v) TFA/water. After washing with 20 ml 0.12% TFA/water, bound material was released by elution with 2 ml of 80% (v/v) acetonitrile/water and concentrated using a Speed-Vac concentrator (Runcorn, UK). The volume was adjusted to 1.0ml with 0.12% (v/v) TFA/water prior to HPLC purification.

HPLC analysis of degraded GIP and Tyr^1-glucitol GIP. Samples were applied to a Vydac C-18 widepore column equilibriated with 0.12% (v/v) TFA/H_2O at a flow rate of 1.0 ml/min. Using 0.1% (v/v) TFA in 70% acetonitrile/H_2O, the concentration of acetonitrile in the eluting solvent was raised from 0% to 31.5% over 15 min, to 38.5% over 30 min and from 38.5% to 70% over 5 min, using linear gradients. The absorbance was monitored at 206 nm and peak areas evaluated using a

model 2221 LKB integrator. Samples recovered manually were concentrated using a Speed-Vac concentrator.

Electrospray ionization mass spectrometry (ESI-MS). Samples for ESI-MS analysis containing intact and degradation fragments of GIP (from DPP IV and plasma incubations) as well as Tyr^1-glucitol GIP, were further purified by HPLC. Peptides were dissolved (approximately 400 pmol) in 100 μl of water and applied to the LCQ benchtop mass spectrometer (Finnigan MAT, Hemel Hempstead, UK) equipped with a microbore C-18 HPLC column (150 x 2.0mm, Phenomenex, UK, Ltd, Macclesfield). Samples (30μl direct loop injection) were injected at a flow rate of 0.2ml/min, under isocratic conditions 35% (v/v) acetonitile/water. Mass spectra were obtained from the quadripole ion trap mass analyzer and recorded. Spectra were collected using full ion scan mode over the mass-to-charge (m/z) range 150-2000. The molecular masses of GIP and related structures were determined from ESI-MS profiles using prominent multiple charged ions and the following equation $M_r = iM_i - iM_h$ (where M_r = molecular mass; M_i = m/z ratio; i = number of charges; M_h = mass of a proton).

***In vivo* biological activity of GIP and Try^1-glucitol GIP.** Effects of GIP and Tyr^1-glucitol GIP on plasma glucose and insulin concentrations were examined using 10-12 week old male Wistar rats. The animals were housed individually in an air conditioned room and 22±2°C with a 12 hour light/12 hour dark cycle. Drinking water and a standard rodent maintenance diet

(Trouw Nutrition, Belfast) were supplied *ad libitum*. Food was withdrawm for an 18 hour period prior to intraperitoneal injection of glucose alone (18mmol/kg body weight) or in combination with either GIP or Tyr^1-glucitol GIP (10 nmol/kg). Test solutions were administered in a final volume of 8 ml/kg body weight. Blood samples were collected at 0, 15, 30 and 60 minutes from the cut tip of the tail of conscious rats into chilled fluoride/heparin microcentrifuge tubes (Sarstedt, Nümbrecht, Germany). Samples were centrifuged using a Beckman microcentrifuge for about 30 seconds at 13,000 g. Plasma samples were aliquoted and stored at -20°C prior to glucose and insulin determinations. All animal studies were done in accordance with the Animals (Scientific Procedures) Act 1986.

Analyses. Plasma glucose was assayed by an automated glucose oxidase procedure using a Beckman Glucose Analyzer II [33]. Plasma insulin was determined by dextran charcoal radioimmunoassay as described previously [34]. Incremental areas under plasma glucose and insulin curves (AUC) were calculated using a computer program (CAREA) employing the trapezoidal rule [35] with baseline subtraction. Results are expressed as mean ± SEM and values were compared using the Student's unpaired *t*-test. Groups of data were considered to be significantly different if $P<0.05$.

Results

Degradation of GIP and Tyr^1-glucitol GIP by DPP IV. Figure 1 illustrates the typical peak profiles obtained from the HPLC separation of the products obtained from the incubation of GIP (Fig 1a) or Tyr^1-glucitol GIP (Fig 1b) with DPP IV for 0, 2, 4 and 12 hours. The retention times of GIP and Tyr^1-glucitol GIP at t=0 were 21.93 minutes and 21.75 minutes respectively. Degradation of GIP was evident after 4 hours incubation (54% intact), and by 12 hours the majority (60% of intact GIP was converted to the single product with a retention time of 21.61 minutes. Tyr^1-glucitol GIP remained almost completely intact throughout 2-12 hours incubation. Separation was on a Vydac C-18 colum using linear gradients of 0% to 31.5% acetonitrile over 15 minutes, to 38.5% over 30 minutes and from 38.5 to 70% acetonitrile over 5 minutes.

Degradation of GIP and Tyr^1-glucitol GIP by human plasma. Figure 2 shows a set of typical HPLC profiles of the products obtained from the incubation of GIP or Tyr^1-glucitol GIP with human plasma for 0 and 4 h. GIP (Fig 2a) with a retention time of 22.06 min was readily metabolised by plasma within 4 hours incubation giving rise to the appearance of a major degradation peak with a retention time of 21.74 minutes. In contrast, the incubation of Tyr^1-glucitol GIP under similar conditions (Fig 2b) did not result in the formation of any detectable degradation fragments during this time with only a single peak being observed with a retention time of 21.77 minutes. Addition of diprotin A, a specific inhibitor of DPP IV, to GIP during the 4 hours incubation completely inhibited degradation of

the peptide by plasma. Peaks corresponding with intact GIP, GIP (3-42) and Tyr^1 - glucitol GIP are indicated. A major peak corresponding to the specific DPP IV inhibitor tripeptide DPA appears in the bottom peanels with retention time of 16-29 min.

Identification of GIP degradation fragments by ESI-MS. Figure 3 shows the monoisotopic molecular masses obtained for GIP, (panel A), Tyr^1-glucitol GIP (panel B) and the major plasma degradation fragment of GIP (panel C) using ESI-MS. The peptides analyzed were purified from plasma incubations as shown in Figure 2. Peptides were dissolved (approximately 400 pmol) in 100µl of water and applied to the LC/MS equipped with a microbore C-18 HPLC column. Samples (30µl direct loop injection) were applied at a flow rate of 0.2ml/min, under isocratic conditions 35% acetonitrile/water. Mass spectra were recorded using a quadripole ion trap mass analyzer. Spectra were collected using full ion scan mode over the mass-to-charge (m/z) range 150-2000. The molecular masses (M_r) of GIP and related structures were determined from ESI-MS profiles using prominent multiple charged ions and the following equation $M_r = iM_i - iM_h$. The exact molecular mass (M_r) of the peptides were calculated using the equation $M_r = iM_i - iM_h$ as defined in Research Design and Methods. After spectral averaging was performed, prominent multiple charges species $(M+3H)^{3+}$ and $(M+4H)^{4+}$ were detected from GIP at m/z 1661.6 and 1246.8, corresponding to intact M_r 4981.8 and 4983.2 Da, respectively (Fig. 3A). Similarly, for Tyr^1-glucitol GIP ($(M+4H)^{4+}$ and $(M+5H)^{5+}$) were detected at m/z 1287.7 and 1030.3, corresponding

to intact molecular masses of M^r 5146.8 and 5146.5 Da, respectively (Fig. 3B). The difference between the observed molecular masses of the quadruply charged GIP and the N-terminally modified GIP species (163.6 Da) indicated that the latter peptide contained a single glucitol adduct corresponding to Tyr^1-glucitol GIP. Figure 3C shows the prominent multiply charged species $(M+3H)^{3+}$ and $(M+4H)^{4+}$ detected from the major fragment of GIP at m/z 1583.8 and 1188.1, corresponding to intact M^r 4748.4 and 4748 Da, respectively (Figure 3C). This corresponds with the theoretical mass of the N-terminally truncated form of the peptide GIP(3-42). This fragment was also the major degradation product of DPP IV incubations (data not shown).

Effects of GIP and Tyr^1-glucitol GIP on plasma glucose homeostasis. Figures 4 and 5 show the effects of intraperitoneal (ip) glucose alone (18mmol.kg) (control group), and glucose in combination with GIP or Tyr^1-glucitol GIP (10nmol/kg) on plasma glucose and insulin concentrations.

(4A) Plasma glucose concentrations after i.p. glucose alone (18mmol/kg) (control group), or glucose in combination with either GIP of Tyr^1-glucitol GIP (10nmol/kg). The time of injection is indicated by the arrow (0 min). **(4B)** Plasma glucose·AUC calues for 0-60 min post injection. Values are mean ± SEM for six rats. $^{**}P<0.01$, $^{***}P<0.001$ compared with GIP and Tyr^1-glucitol GIP; $^{\dagger}P<0.05$, $^{\dagger\dagger}P<0.01$ compared with non-glucated GIP.

(5A). Plasma insulin concentrates after i.p. glucose along (18 mmol/kg) (control group), or glucose in combination with either with GIP or glycated GIP (10nmol/kg). The time of injection is indicated by the arrow. **(5B)** Plasma insulin AUC values were calculated for each of the 3 groups up to 90 minutes post injection. The time of injection is indicated by the arrow (0 min). Plasma insulin AUC values for 0-60 min post injection. Values are mean ± SEM for six rats. **P*<0.05, ***P*<0.001 compared with GIP and Tyr^1-glucitol GIP; †*P*<0.05, ††*P*<0.01 compared with non-glycated GIP.

Compared with the control group, plasma glucose concentrations and area under the curve (AUC) were significantly lower following administration of either GIP or Tyr^1- glucitol GIP (Figure 4A, B). Furthermore, individual values at 15 and 30 minutes together with AUC were significantly lower following administration of Tyr^1-glucitol GIP as compared to GIP. Consistent with the established insulin-releasing properties of GIP, plasma insulin concentrations of both peptide-treated groups were significantly raised at 15 and 30 minutes compared with the values after administration of glucose alone (Figure 5A). The overall insulin responses, estimated as AUC were also signigicantly greater for the two peptide-treated groups (Figure 5B). Despite lower prevailing glucose concentrations than the GIP-treated group, plasma insulin response, calculated as AUC, following Tyr^1-glucitol GIP was significantly greater than after GIP (Figure 5B). The significant elevation of plasma insulin at 30 minutes is of particular note, suggesting that the insulin-

releasing action of Tyr^1-glucitol GIP is more protracted than GIP even in the face of a diminished glycemic stimulus (Figures 4A, 5A).

Discussion

The forty two amino acid GIP is an important incretin hormone released into the circulation from endocrine K-cells of the duodenum and jejunum following ingestion of food . The high degree of structural conservation of GIP among species supports the view that this peptide plays and important role in metabolism. Secretion of GIP is stimulateed directly by actively transported nutrients in the gut lumen without a notable input from autonomic nerves. The most important stimulants of GIP release are simple sugars and unsaturated long chain fatty acids, with amino acids exerting weaker effects. As with tGLP-1, the insulin-releasing effect of GIP is strictly glucose-dependent. This affords protection against hypoglycemia and thereby fulfils one of the most desirable features of any current or potentially new antidiabetic drug.

The present results demonstrate for the first time that Tyr^1-glucitol GIP displays profound resistance to serum and DPP IV degradation. Using ESI-MS the present study showed that native GIP was rapidly cleaved *in vitro* to a major 4748.4 Da degradation product, corresponding to GIP(3-42) which confirmed previous findings using matrix-assisted laser desorption ionization time-of-flight mass spectrometry. Serum degradation was

completely inhibited by diprotin A (Ile-Pro-Ile), a specific competitive inhibitor of DPP IV, confirming this as the main enzyme for GIP inactivation *in vivo*. In contrast, Tyr^1-glucitol GIP remained almost completely intact after incubation with serum or DPP IV for up to 12 hours. This indicates that glycation of GIP at the amino-terminal Tyr^1 residue masks the potential cleavage site from DPP IV and prevents removal of the Tyr^1-Ala^2 dipeptide from the N-terminus preventing the formation of GIP(3-42).

Consistent with *in vitro* protection against DPP IV, administration of Tyr^1-glucitol GIP significantly enhanced the antihyperglycemic activity and insulin-releasing action of the peptide when administered with glucose to rats. Native GIP enhanced insulin release and reduced the glycemic excursion as observed in many previous studies. However, amino-terminal glycation of GIP increased the insulin-releasing and antihyperglycemic actions of the peptide by 62% and 38% respectively, as estimated from AUC measurements. Detailed kinetic analysis is difficult due to necessary limitation of sampling times, but the greater insulin concentrations following Tyr^1-glucitol GIP as opposed to GIP at 30 minutes post-injection is indicative of a longer half-life. The glycemic rise was modest in both peptide-treated groups and glucose concentrations following injection of Tyr^1-glucitol GIP were consistently lower than after GIP. Since the insulinotropic actions of GIP are glucose-dependent, it is likely that the relative insulin-releasing potency

of Tyr^1-glucitol GIP is greatly underestimated in the present *in vivo* experiments.

In vitro studies in the laboratory of the present inventors using glucose-responsive clonal B-cells showed that the insulin-releasing potency of Tyr^1-glucitol GIP was several order of magnitude greater than GIP and that its effectiveness was more sensitive to change of glucose concentrations within the physiological range. Together with the present *in vivo* observations, this suggests that N-terminal glycation of GIP confers resistance to DPP IV degradation whilst enhancing receptor binding and insulin secretory effects on the B-cell. These attributes of Tyr^1-glucitol GIP are fully expressed *in vivo* where DPP IV resistance impedes degradation of the peptide to GIP(3-42), thereby prolonging the half-life and ehancing effective concentrations of the intact biologically active peptide. It is thus possible that glycated GIP is enhancing insulin secretion *in vivo* both by enhanced potency at the receptor as well as improving DPP IV resistance. Thus numerous studies have shown that GIP (3-42) and other N-terminally modified fragments, including GIP(4-42), and GIP (17-42) are either weakly effective or inactive in stimulating insulin release. Furthermore, evidence exists that N-terminal deletions of GIP result in receptor antagonist properties in GIP receptor transfected Chinese hamster kidney cells [9], suggesting that inhibition of GIP catabolism would also reduce the possible feedback antagonism at the receptor level by the truncated GIP(3-42).

In addition to its insulinotopic actions, a number of other potentially important extrapancreatic actions of GIP may contribute to the enhanced antihyperglycemic activity and other beneficial metabolic effects of Tyr^1-glucitol GIP. These include the stimulation of glucose uptake in adipocytes, increased synthesis of fatty acids and activation of lipoprotein lipase in adipose tissue. GIP also promotes plasma triglyceride clearance in response to oral fat loading. In liver, GIP has been shown to enhance insulin-dependent inhibition of glycogenolysis. GIP also reduces both glucagon-stimulated lipolysis in adipose tissue as well as hepatic glucose production. Finally, recent findings indicate that GIP has a potent effect on glucose uptake and metabolism in mouse isolated diaphragm muscle. This latter action may be shared with tGLP-1 and both peptides have additional benfits of stimulating somatostatin secretion and slowing down gastric emptying and nutrient absorption.

In conclusion, this study has demonstrated for the first time that the glycation of GIP at the amino-terminal Tyr^1 residue limits GIP catabolism through impairment of the proteolytic actions of serum petidases and thus prolongs its half-life *in vivo*. This effect is accompanied by enhanced antihyperglycemic activity and raised insulin concentrations *in vivo*, suggesting that such DPP IV resistant analogues should be explored alongside tGLP-1 as potentially useful therapeutic agents for NIDDM. Tyr^1-glucitol GIP appears to be particularly interesting in this regard since such amino-terminal

modification of GIP enhances rather than impairs glucose-dependent insulinotropic potency as was observed recently for tGLP-1.

Example 3

This example further looked at the ability of additional N-terminal structural modifications of GIP in preventing inactivation by DPP and in plasma and their associated increase in both the insulin-releasing potency and potential therapeutic value. Native human GIP, glycated GIP, acetylated GIP and a number of GIP analogues with N-terminal amino acid substitutions were tested.

Materials and Methods

Reagents

High-performance liquid chromatography (HPLC) grade acetonitrile was obtained from Rathburn (Walkersburn, Scotland). Sequencing grade trifluoroacetic acid (TFA) was obtained from Aldrich (Poole, Dorset, UK). Dipeptidyl peptidase IV was purchased from Sigma (Poole, Dorset, UK), and Diprotin A was purchased from Calbiochem Novabiochem (Beeston, Nottingham, UK). RPMI 1640 tissue culture medium, foetal calf serum, penicillin and streptomycin were all purchased from Gibco (Paisley, Strathclyde, UK). All water used in these experiments was purified using a Milli-Q, Water Purification System (Millipore, Millford, MA, USA).

All other chemicals used were of the highest purity available.

Synthesis of GIP and N-terminally modified GIP analogues

GIP, GIP(Abu2), GIP(Sar2), GIP(Ser2), GIP(Gly2) and GIP(Pro3) were sequentially synthesised on an Applied Biosystems automated peptide synthesizer (model 432A) using standard solid-phase Fmoc procedure, starting with an Fmoc-Gln-Wang resin. Following cleavage from the resin by trifluoroacetic acid: water, thioanisole, ethanedithiol (90/2.5/5/2.5, a total volume of 20 ml/g resin), the resin was removed by filtration and the filtrate volume was decreased under reduced pressure. Dry diethyl ether was slowly added until a precipitate was observed. The precipitate was collected by low-speed centrifugation, resuspended in diethyl ether and centrifuged again, this procedure being carried out at least five times. The pellets were then dried in vacuo and judged pure by reversed-phase HPLC on a Waters Millennium 2010 chromatography system (Software version 2.1.5.). N-terminal glycated and acetylated GIP were prepared by minor modification of a published method.

Electrospray ionization-mass spectrometry (ESI-MS) was carred out as described in Example 2.

Degradation of GIP and novel GIP analogues by DPP IV and human plasma was carried out as described in Example 2.

Culture of insulin secreting cells

BRIN-BD11 cells [30] were cultured in sterile tissue culture flasks (Corning, Glass Works, UK) using RPMI-1640 tissue culture medium containing 10% (v/v) foetal calf serum, 1% (v/v) antibiotics (100 U/ml penicillin, 0.1 mg/ml streptomycin) and 11.1 mM glucose. The cells were maintained at 37°C in an atmosphere of 5% CO_2 and 95% air using a LEEC incubator (Laboratory Technical Engineering, Nottingham, UK).

Acute tests for insulin secretion

Before experimentation, the cells were harvested from the surface of the tissue culture flasks with the aid of trypsin/EDTA (Gibco), seeded into 24-multiwell plates (Nunc, Roskilde, Denmark) at a density of 1.5 x 105 cells per well, and allowed to attach overnight at 37°C. Acute tests for insulin release were preceded by 40 min pre-incubation at 37°C in 1.0 ml Krebs Ringer bicarbonate buffer (115 mM NaCl, 4.7 mM KCl, 1.28 mM $CaCl_2$, 1.2 mM KH_2PO_4, 1.2 mM $MgSO_4$, 10 mM $NaHCO_3$, 5 g/l bovine serum albumin, pH 7.4) supplemented with 1.1 mM glucose. Test incubations were performed (n=12) at two glucose concentrations (5.6 mM and 16.7 mM) with a range of concentrations (10^{-13} to 10^{-8} M) of GIP or GIP analogues. After 20 min incubation, the buffer was removed from each well and aliquots (200 μl) were used for measurement of insulin by radioimmunoassay [31].

Statistical analysis

[redacted]

Results are expressed as mean ± S.E.M. and values were compared using the Student's unpaired t-test. Groups of data were considered to be significantly different if $P < 0.05$.

Results and Discussion

Structural identification of GIP and GIP analogues by ESI-MS

The monoisotopic molecular masses of the peptides were determined using ESI-MS. After spectral averaging was performed, prominent multiple charged species $(M+3H)^{3+}$ and $(M+4H)^{4+}$ were detected for each peptide. Calculated molecular masses confirmed the structural identity of synthetic GIP and each of the N-terminal analogues.

Degradation of GIP and novel GIP analogues by DPP-IV

Figs. 6-11 illustrate the typical peak profiles obtained from the HPLC separation of the reaction products obtained from the incubation of GIP, GIP(Abu2), GIP(Sar2), GIP(Ser2), glycated GIP and acetylated GIP with DPP IV, for 0, 2, 4, 8 and 24 h. The results summarised in Table 1 indicate that glycated GIP, acetylated GIP, GIP(Ser2) are GIP(Abu2) more resistant than native GIP to in vitro degradation with DPP IV. From these data GIP(Sar2) appears to be less resistant.

Degradation of GIP and GIP analogues by human plasma

Figs. 12-16 show a representative set of HPLC profiles obtained from the incubation of GIP and GIP analogues with human plasma for 0, 2, 4, 8 and 24 h. Observations were also made after incubation for 24 h in the presence of DPA. These results are summarised in Table 2 are broadly comparable with DPP IV incubations, but conditions which more closely mirror in vivo conditions are less enzymatically severe. GIP was rapidly degraded by plasma. In comparison, all analogues tested exhibited resistance to plasma degradation, including GIP(Sar2) which from DPP IV data appeared least resistant of the peptides tested. DPA substantially inhibited degradation of GIP and all analogues tested with complete abolition of degradation in the cases of GIP(Abu2), GIP(Ser2) and glycated GIP. This indicates that DPP IV is a key factor in the in vivo degradation of GIP.

Dose-dependent effects of GIP and novel GIP analogues on insulin secretion

Figs. 17-30 show the effects of a range of concentrations of GIP, GIP(Abu2), GIP(Sar2), GIP(Ser2), acetylated GIP, glycated GIP, GIP(Gly2) and GIP(Pro3) on insulin secretion from BRIN-BD11 cells at 5.6 and 16.7 mM glucose. Native GIP provoked a prominent and dose-related stimulation of insulin secretion. Consistent with previous studies [28], the glycated GIP analogue exhibited a considerably greater insulinotropic response compared with native peptide. N-terminal acetylated GIP exhibited a similar pattern and the GIP(Ser2) analogue also evoked a strong

response. From these tests, GIP(Gly2) and GIP(Pro3) appeared to the least potent analogues in terms of insulin release. Other stable analogues tested, namely GIP(Abu2) and GIP(Sar2), exhibited a complex pattern of responsiveness dependent on glucose concentration and dose employed. Thus very low concentrations were extremely potent under hyperglycaemic conditions (16.7 mM glucose). This suggests that even these analogues may prove therapeutically useful in the treatment of type 2 diabetes where insulinotropic capacity combined with in vivo degradation dictates peptide potency.

Table 1 : % Intact peptide remaining after incubation with DPPIV

Peptide	% Intact peptide remaining after time (h)				
	0	2	4	8	24
GIP 1-42	100	52 ± 1	23 ± 1	0	0
Glycated GIP	100	100	100	100	100
GIP (Abu^2)	100	38 ± 1	28 ± 2	0	0
GIP (Ser^2)	100	77 ± 2	60 ± 1	32 ± 4	0
GIP (Sar^2)	100	28 ± 2	8	0	0
N-Acetyl-GIP	100	100	100	100	0

Table 2 : % Intact peptide remaining after incubation with human plasma

Peptide	% Intact peptide remaining after incubations with human plasma					
	0	2	4	8	24	DPA
GIP 1-42	100	52 ± 1	23 ± 1	0	0	68 ± 2
Glycated GIP	100	100	100	100	100	100
GIP (Abu^2)	100	38 ± 1	28 ± 2	0	0	100
GIP (Ser^2)	100	77 ± 2	60 ± 1	32 ± 4	0	63 ± 3
GIP (Sar^2)	100	28 ± 2	8	0	0	100

Tables represent the percentage of intact peptide (i.e. GIP 1-42) relative to the major degradation product GIP 3-42. Values were taken from HPLC traces performed in triplicate and the mean and S.E.M. values calculated. DPA is diprotin A, a specific inhibitor of DPPIV.

CLAIMS

1. A peptide analogue of GIP (1-42) comprising at least 15 amino acid residues from the N terminal end of GIP (1-42) having a least one amino acid substitution or modification at position 1-3 and not including Tyr^1 glucitol GIP (1-42).

2. A peptide analogue as claimed in claim 1 including modification by fatty acid addition at an epsilon amino group of at least one lysine residue.

3. A peptide analogue of biologically active GIP (1-42) wherein the analogue is Tyr^1 glucitol GIP (1-42) modified by fatty acid addition at an epsilon amino group of at least one lysine residue.

4. A peptide analogue as claimed in any of the preceding claims wherein the substitution or modification is chosen from the group comprising D-amino acid substitutions in 1, 2 and/or 3 positions and/or N terminal glycation, alkylation, acetylation or acylation.

5. A peptide analogue as claimed in any of the preceding claims wherein the amino acid in the 2 or 3 position is substituted by lysine, serine, 4-amino butyric, Aib, D-alanine, Sarcosine or Proline.

6. An analogue as claimed in any of the preceding claims wherein the N terminus is modified by one

of the group of modifications include glycation, alkylation, acetylation or by the addition of an isopropyl group.

7. Use of an analogue as claimed in any of the preceding claims in the preparation of a medicament for the treatment of diabetes.

8. A pharmaceutical composition including an analogue as claimed in any of the preceding claims.

9. A pharmaceutical composition as claimed in claim 8 in admixture with a pharmaceutically acceptable excipient.

10. A method of N-terminally modifying GIP or analogues thereof the method comprising the steps of synthesising the peptide from the C terminal to the penultimate N terminal amino acid, adding tyrosine as a F-moc protected Tyr(tBu)-Wang resin, deprotecting the N-terminus of the tyrosine and reacting with modifying agent, allowing the reaction to proceed to completion, cleaving the modified tyrosine from the Wang resin and adding the modified tyrosine to the peptide synthesis reaction.

11. A method as claimed in claim 10 wherein the modifying agent is chosen from the group comprising glucose, acetic anhydride or pyroglutamic acid.

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(54) Title: ANALOGS OF GASTRIC INHIBITORY PEPTIDE AND THEIR USE FOR TREATMENT OF DIABETES

(57) Abstract: The present invention provides peptides which stimulate the release of insulin. The peptides, based on GIP 1-42 include substitutions and/or modifications which enhance and influence secretion and/or have enhanced resistance to degradation. The invention also provides a process of N terminally modifying GIP and the use of the peptide analogues for treatment of diabetes.

Fig. 1a

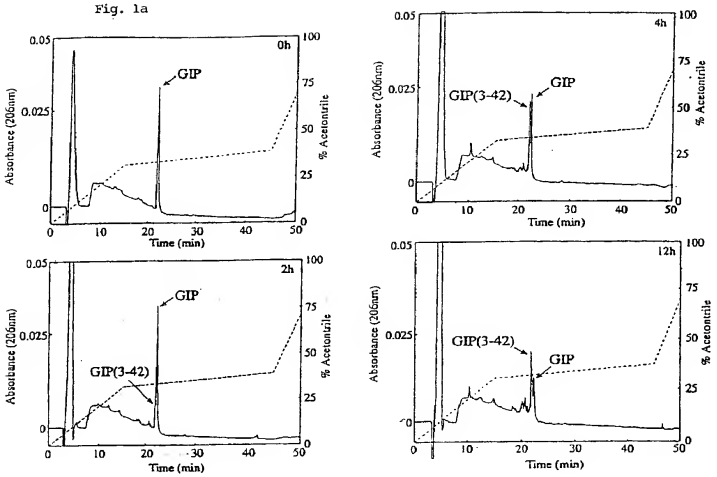
0h
GIP
2h
GIP
GIP(3-42)
4h
GIP(3-42)
GIP
12h
GIP(3-42)
GIP
Absorbance (206nm)
% Acetonitrile
Time (min)

Fig. 1b

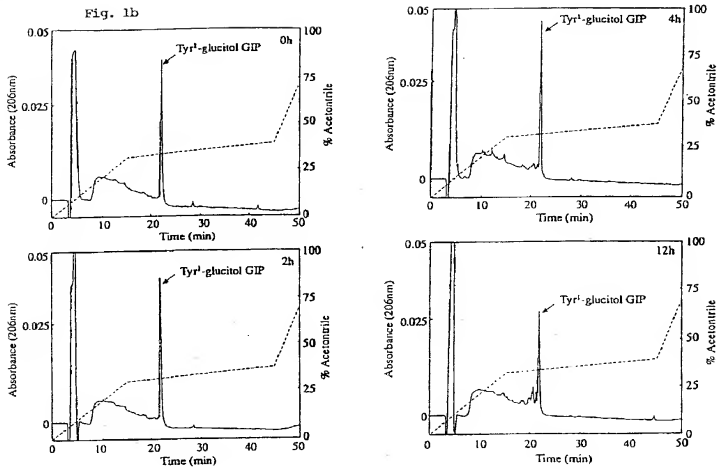
Tyr¹-glucitol GIP
0h
2h
4h
12h
Absorbance (206nm)
% Acetonitrile
Time (min)

Fig. 2a

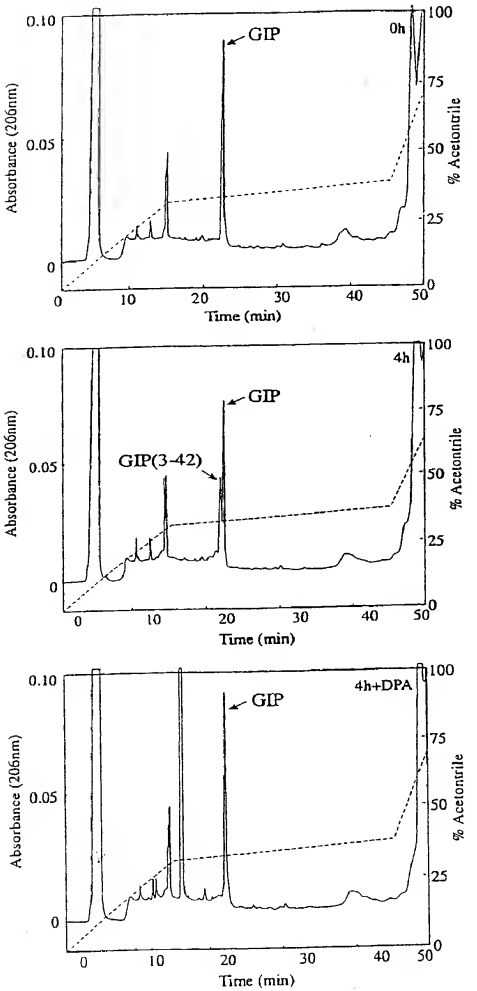
0.10
0.05
0
Absorbance (206nm)
GIP
0h
100
75
50
25
0
% Acetontrile
0
10
20
30
40
50
Time (min)
4h
GIP(3-42)
4h+DPA

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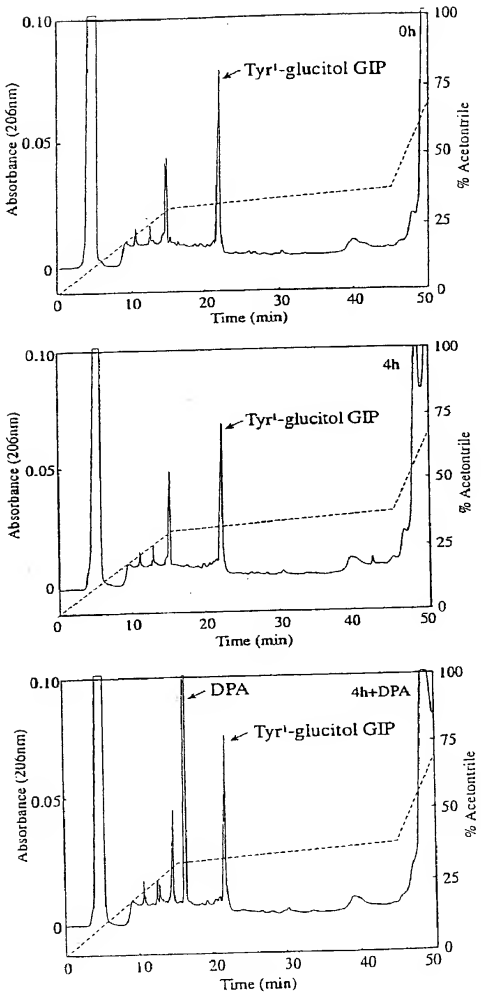
Fig. 2b



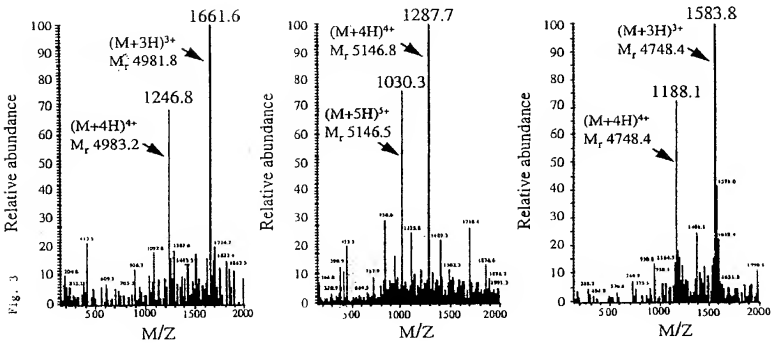


Fig. 3

Fig. 4

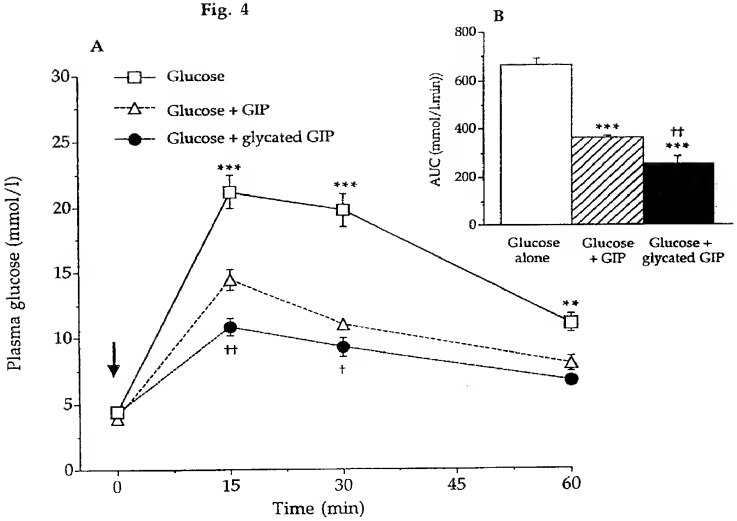
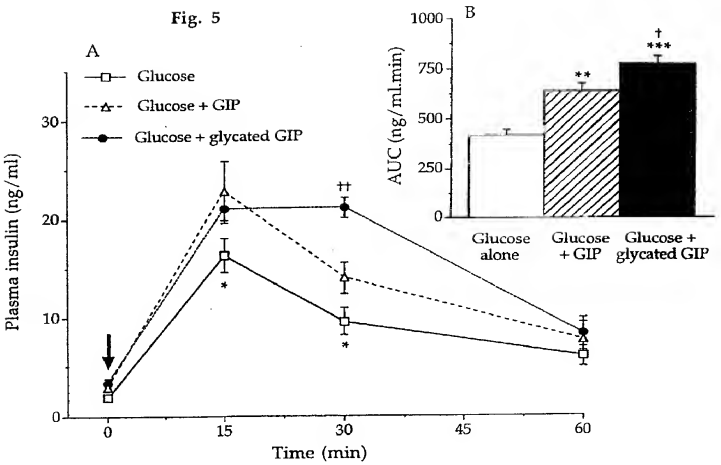

A
Glucose
Glucose + GIP
Glucose + glycated GIP
Plasma glucose (mmol/l)
Time (min)
B
AUC (mmol/l.min)
Glucose alone
Glucose + GIP
Glucose + glycated GIP

Fig. 5


A
Glucose
Glucose + GIP
Glucose + glycated GIP
Plasma insulin (ng/ml)
30
20
10
0
0
15
30
45
60
Time (min)
B
AUC (ng/ml.min)
1000
750
500
250
0
Glucose alone
Glucose + GIP
Glucose + glycated GIP

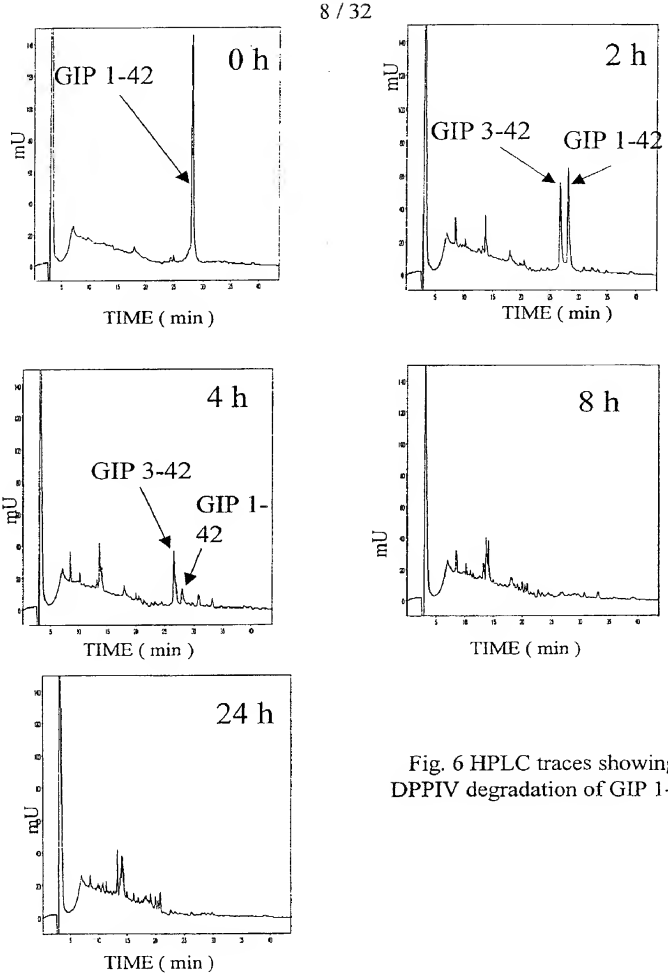


Fig. 6 HPLC traces showing DPPIV degradation of GIP 1-42

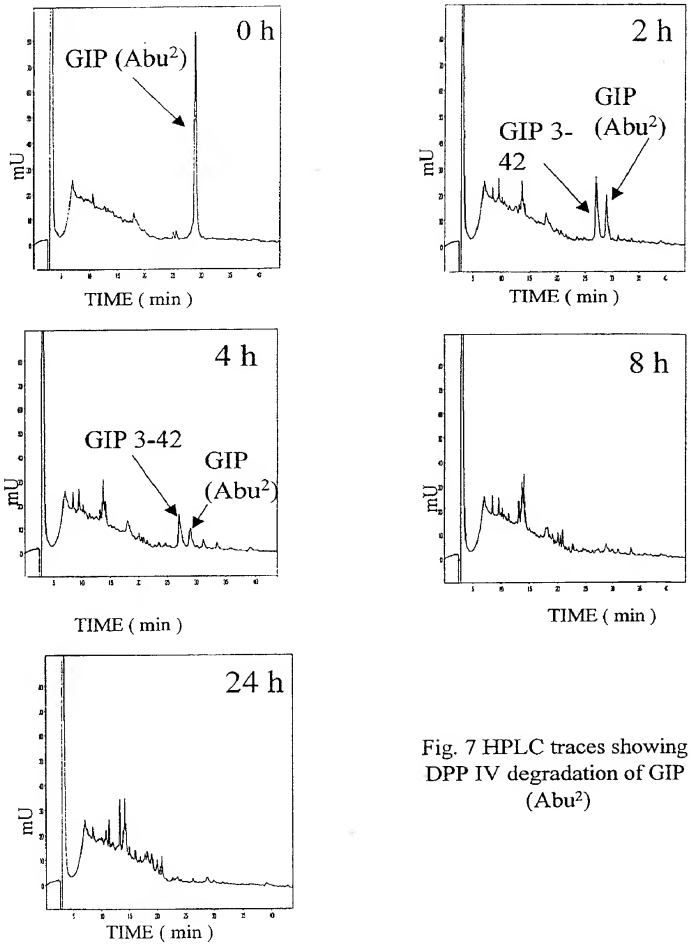


Fig. 7 HPLC traces showing DPP IV degradation of GIP (Abu²)

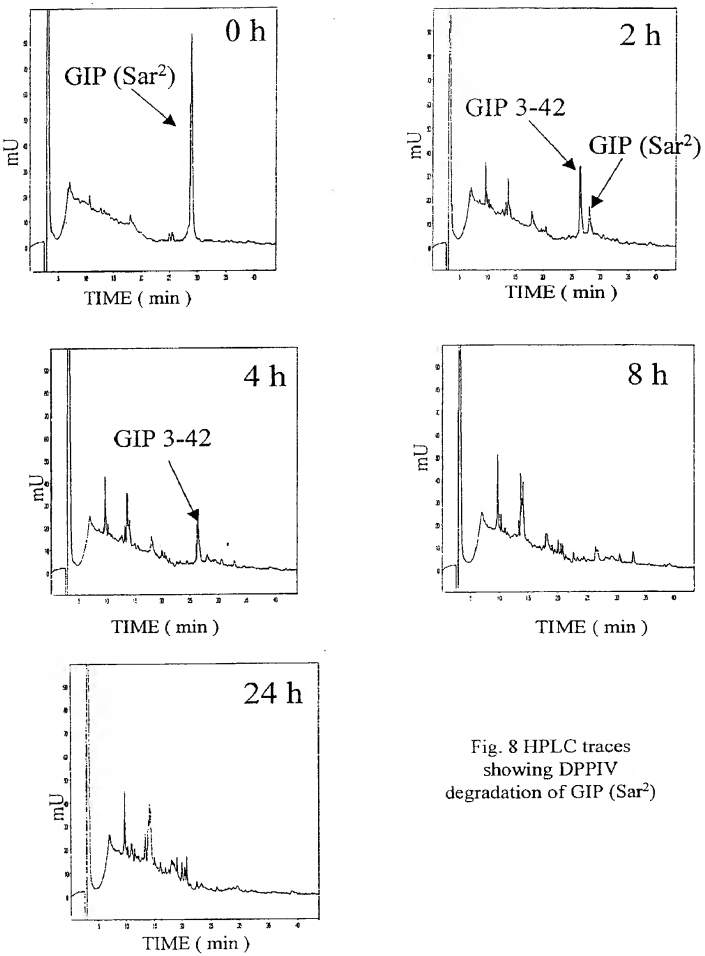


Fig. 8 HPLC traces showing DPPIV degradation of GIP (Sar^2)

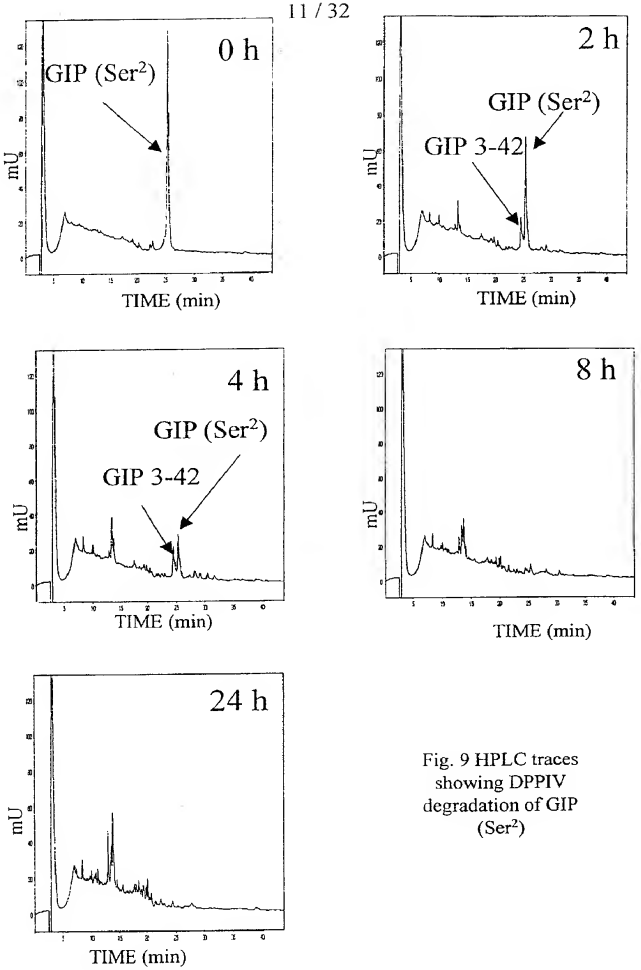


Fig. 9 HPLC traces showing DPPIV degradation of GIP (Ser^2)

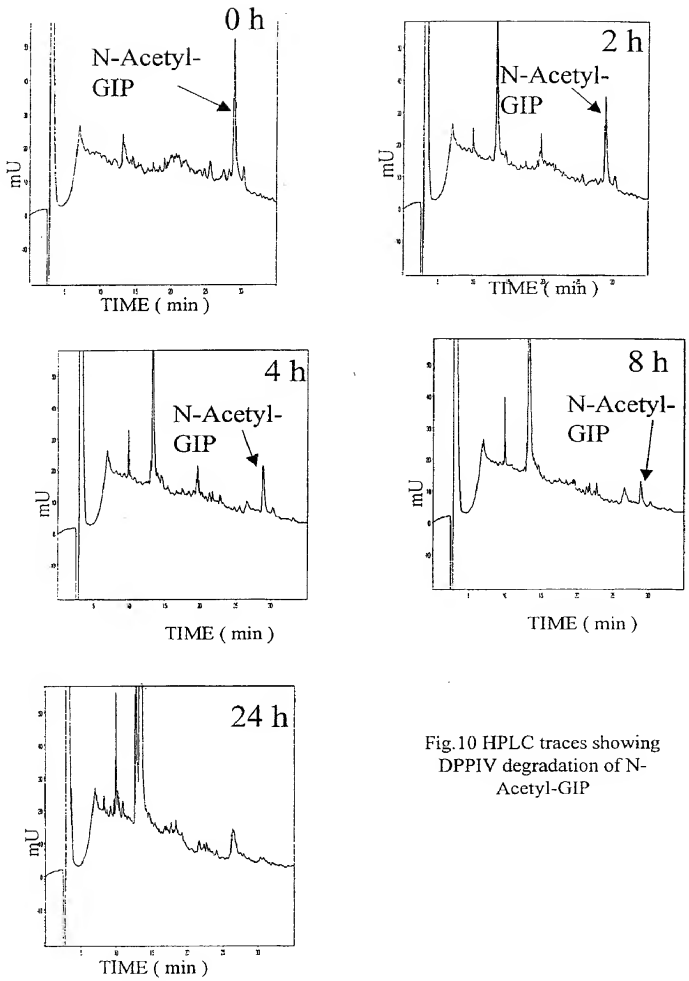


Fig.10 HPLC traces showing DPPIV degradation of N-Acetyl-GIP

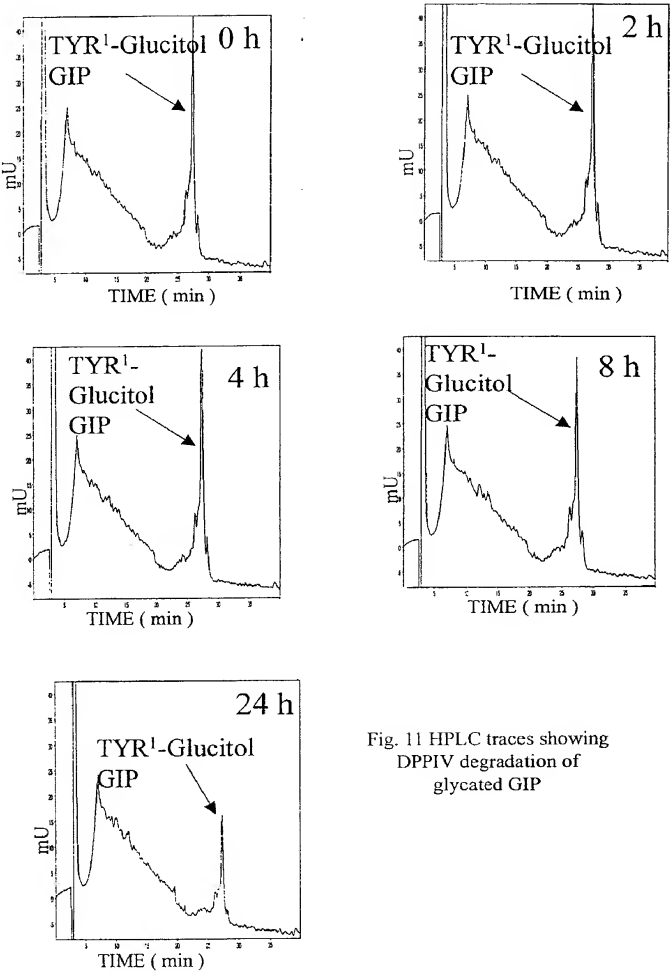


Fig. 11 HPLC traces showing DPPIV degradation of glycated GIP

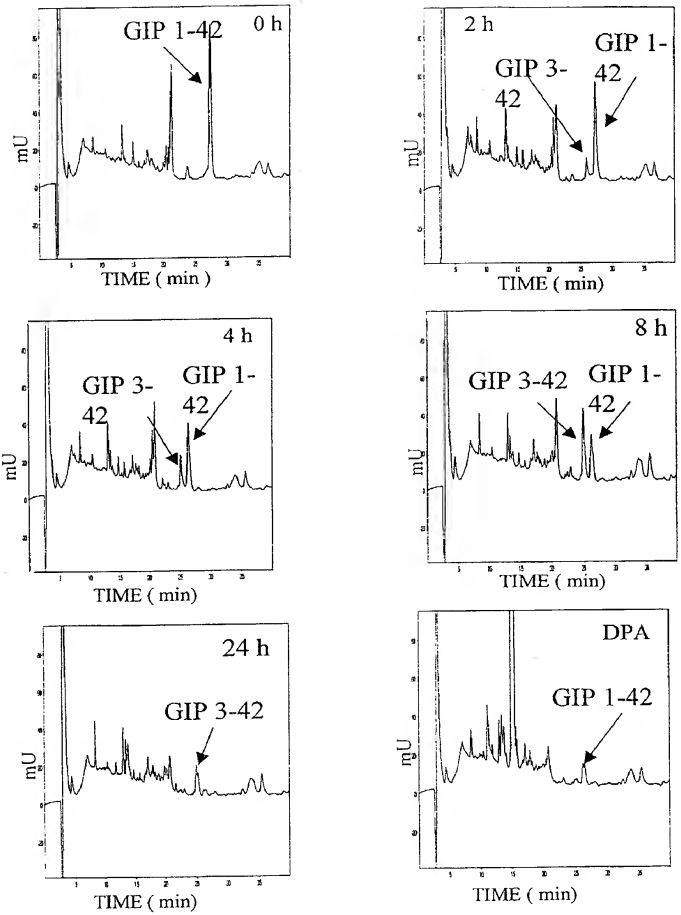


Fig.12. HPLC traces showing human plasma degradation of GIP

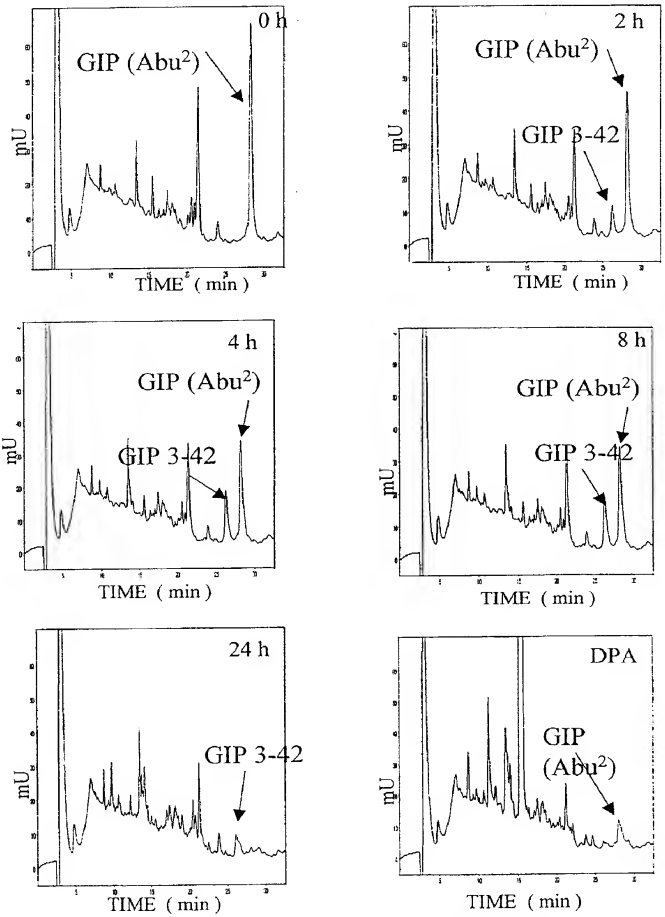


Fig. 13. HPLC traces showing human plasma degradation of GIP (Abu²)

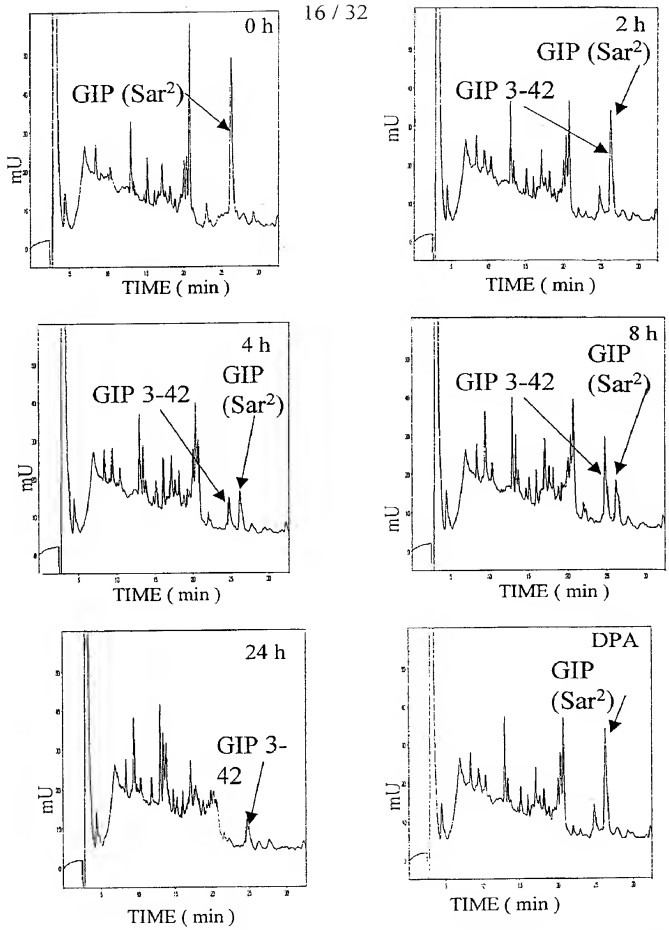


Fig. 14. HPLC traces showing human plasma degradation of GIP (Sar^2)

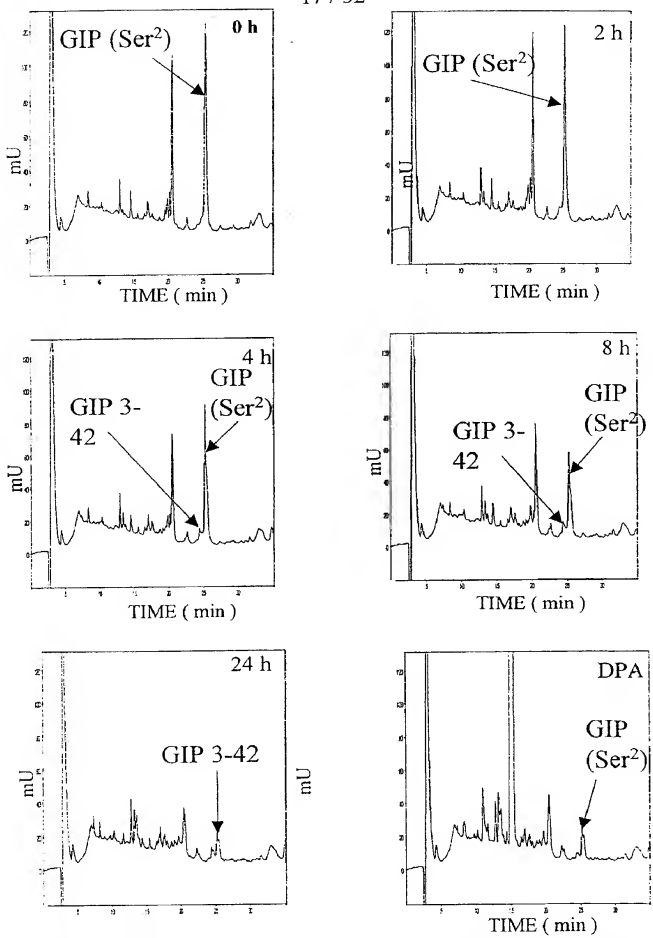


Fig. 15 HPLC traces showing human plasma degradation of GIP(Ser 2)

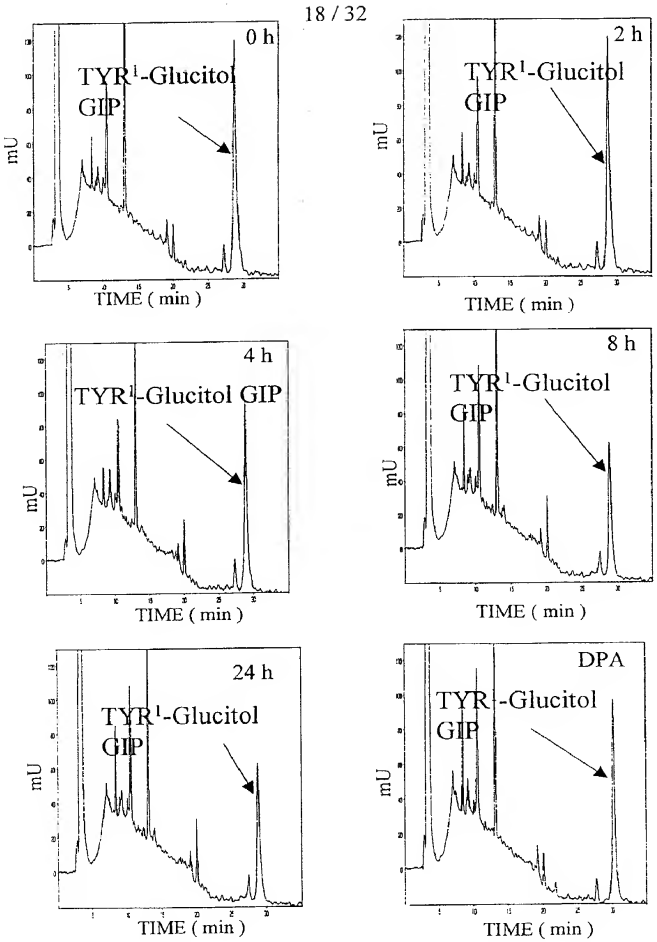
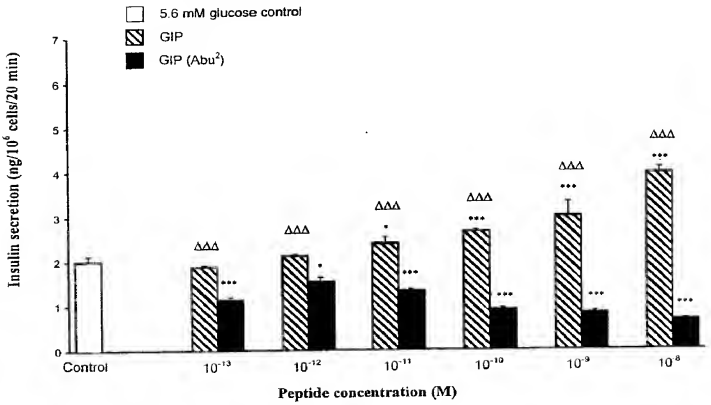


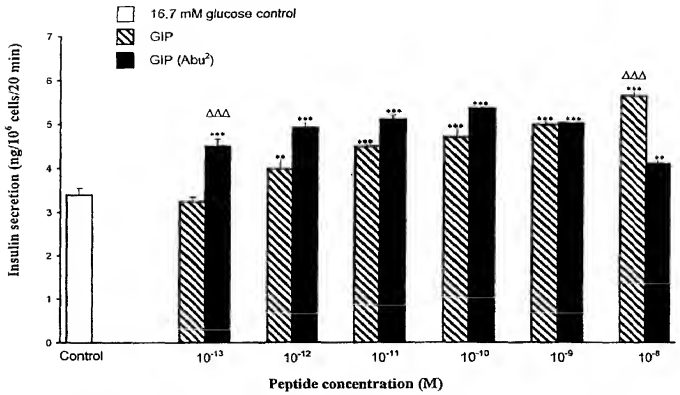
Fig. 16. HPLC traces showing human plasma degradation of glycated GIP

Fig.17. Graph showing the effects of various concentrations of GIP and GIP (Abu^2) on insulin release from BRIN-BD11 cells incubated at 5.6 mM glucose



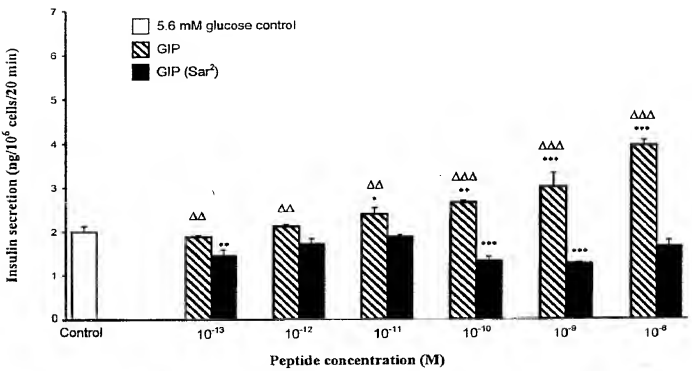
Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (5.6mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Abu^2) at the same concentration.

Fig. 18. Graph showing the effects of various concentrations of GIP and GIP (Abu^2) on insulin release from BRIN-BD11 cells incubated at 16.7 mM glucose.



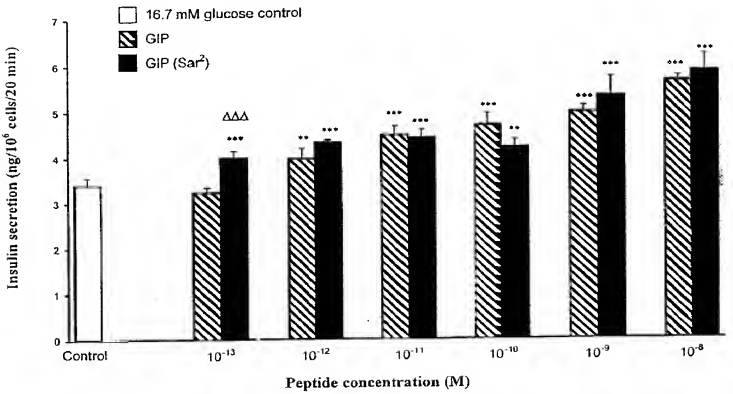
Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (16.7 mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Abu^2) at the same concentration.

Fig.19. Graph showing the effects of various concentrations of GIP and GIP (Sar^2) on insulin release from BRIN-BD11 cells incubated at 5.6 mM glucose



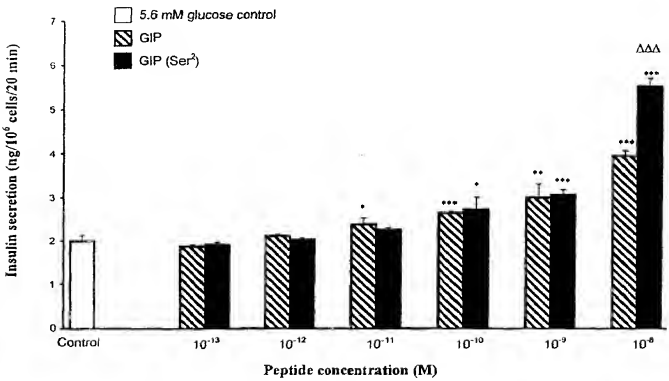
Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (5.6mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Sar^2) at the same concentration.

Fig. 20. Graph showing the effects of various concentrations of GIP and GIP (Sar^2) on insulin release from BRIN-BD11 cells incubated at 16.7 mM glucose



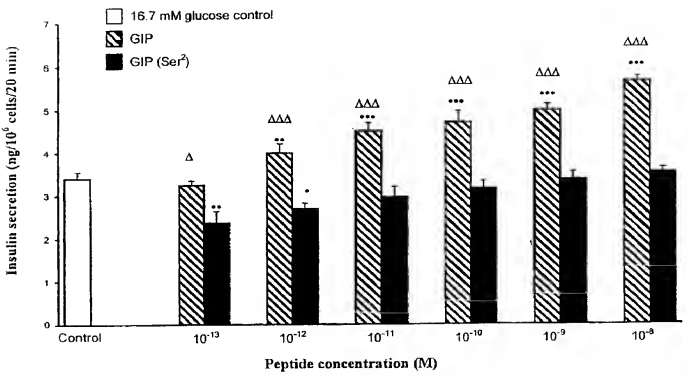
Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (16.7 mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Sar^2) at the same concentration.

Fig.21. *Graph showing the* effects of various concentrations of GIP and GIP (Ser^2) on insulin release from BRIN-BD11 cells incubated at 5.6 mM glucose



Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (5.6mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Ser^2) at the same concentration.

Fig. 22. Graph showing the effects of various concentrations of GIP and GIP (Ser^2) on insulin release from BRIN-BD11 cells incubated at 16.7 mM glucose



Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (16.7 mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Ser^2) at the same concentration.

Fig. 23 Graph showing the effects of various concentrations of GIP 1-42 and N-Acetyl-GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 5.6 mM glucose

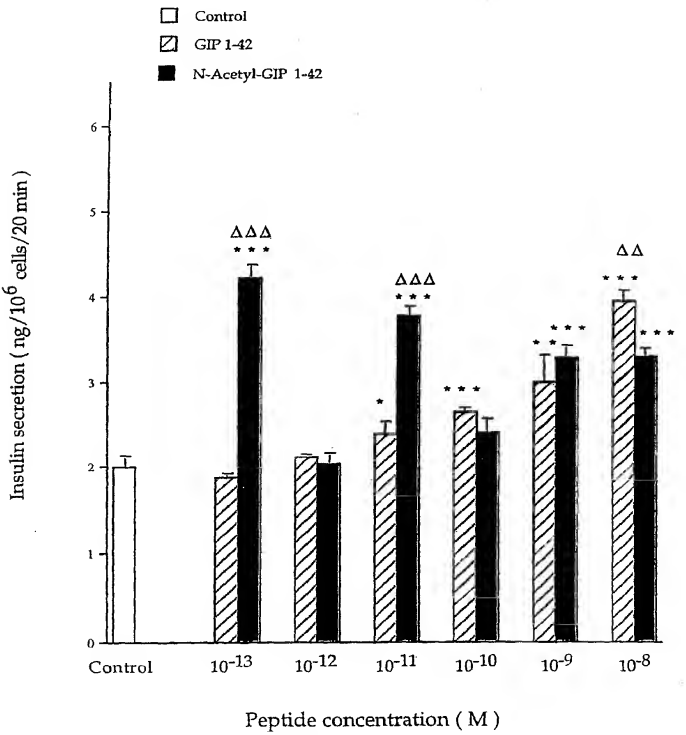


Fig. 24 Graph showing the effects of various concentrations of GIP 1-42 and N-Acetyl-GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 16.7 mM glucose

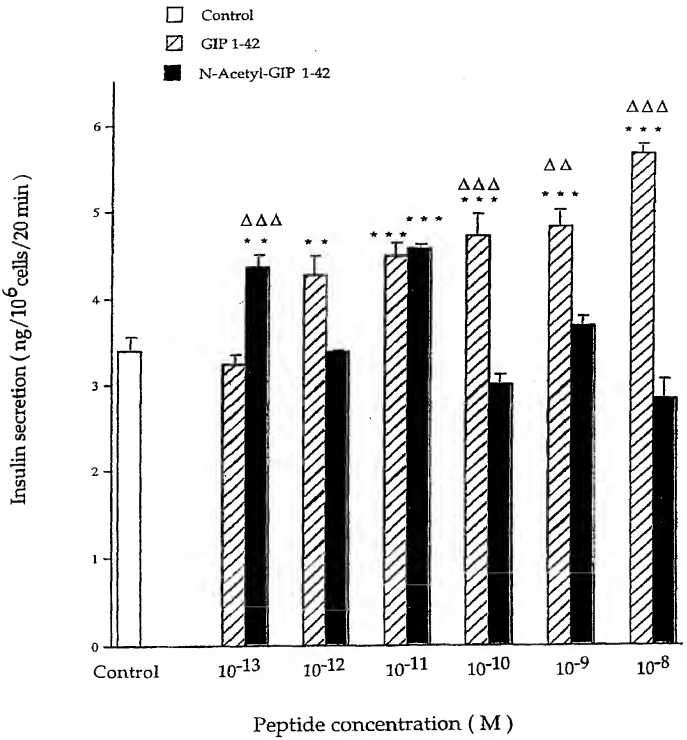


Fig. 25 Graph showing the effects of various concentrations of GIP 1-42 and glycated GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 5.6 mM glucose

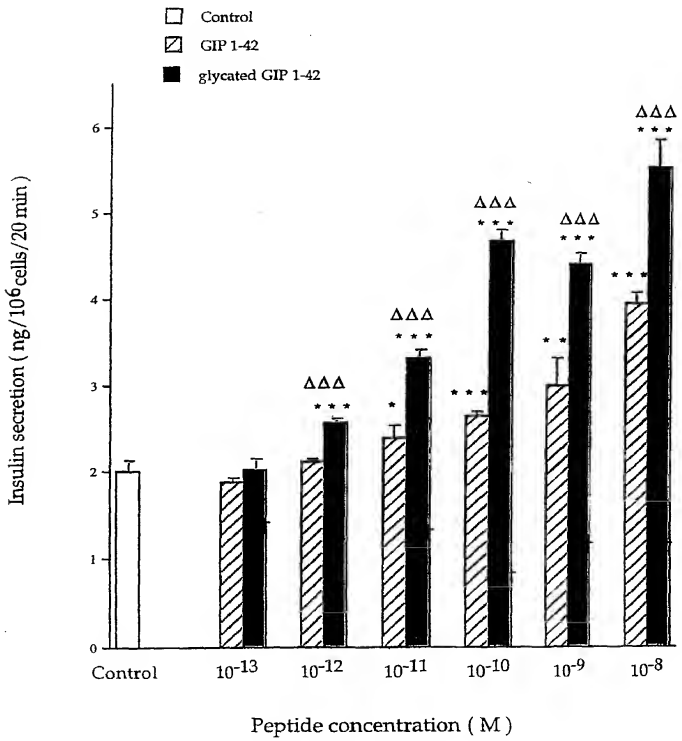


Fig. 26 Graph showing the effects of various concentrations of GIP 1-42 and glycated GIP 1-42 on insulin release from BRIN-BD11 cells incubated at 16.7 mM glucose

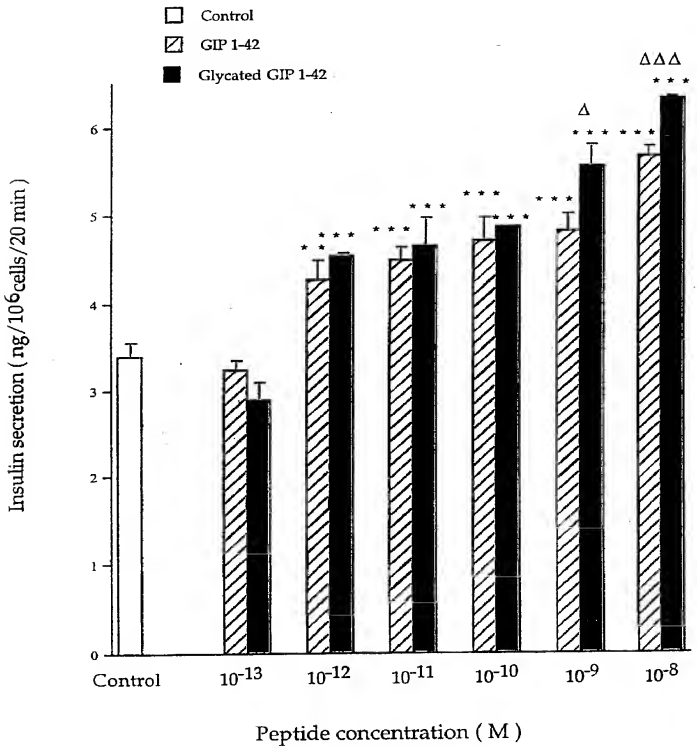
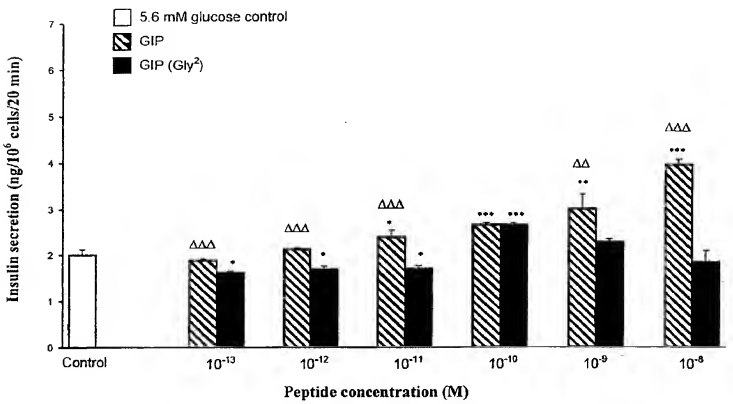
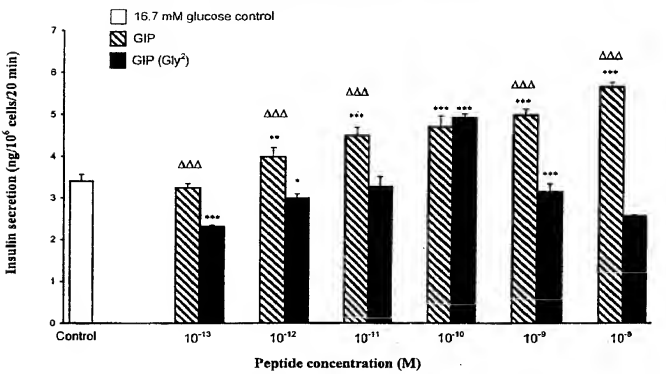


Fig. 27 Graph showing the effects of various concentrations of GIP and GIP (Gly^2) on insulin release from BRIN-BD11 cells incubated at 5.6 mM glucose



Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (5.6mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Gly^2) at the same concentration.

Fig. 28 Graph showing the effects of various concentrations of GIP and GIP (Gly^2) on insulin release from BRIN-BD11 cells incubated at 16.7 mM glucose



Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (16.7 mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Gly^2) at the same concentration.

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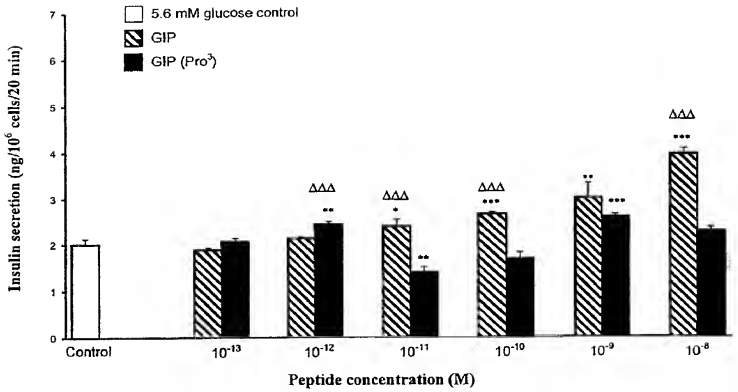
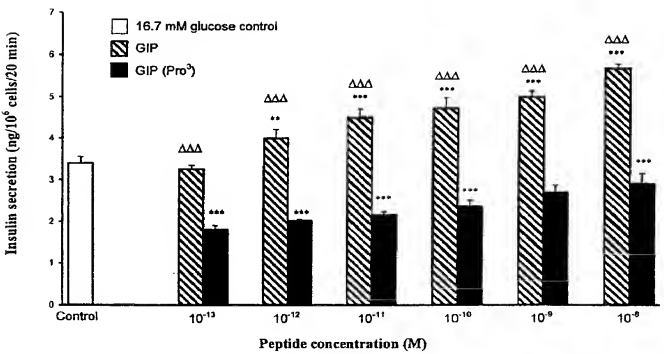
Fig. 29 Graph showing the effects of various concentrations of GIP and GIP (Pro^3) on insulin release from BRIN-BD11 cells incubated at 5.6 mM glucose



Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (5.6mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Pro^3) at the same concentration.

Fig. 30 Graph showing the effects of various concentrations of GIP and GIP (Pro^3) on insulin release from BRIN-BD11 cells incubated at 16.7 mM glucose



Values are means ± S.E.M. for 12 separate observations. $^{*}P< 0.05$, $^{**}P< 0.01$, $^{***}P<0.001$ compared to control (16.7 mM glucose alone). $^{\Delta}P<0.05$, $^{\Delta\Delta}P<0.01$, $^{\Delta\Delta\Delta}P<0.001$ compared to GIP (Pro^3) at the same concentration.

PATENT
Attorney Docket No. 8830-8

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are stated below next to my name:

I believe I am the original, first, and sole inventor (if only one name is listed below) or an original, first, and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

PEPTIDE

---

the specification of which is attached hereto unless the following box is checked

☒ was filed on March 29, 2000 as Application No. _____ or PCT Application No. PCT/GB00/01089 and amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with 37 CFR §1.56.

I hereby claim foreign priority benefits under 35 U.S.C. §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate, or §365(a) of any PCT international application which designated at least one country other than the United States, listed below and have also identified below any foreign application for patent or inventor's certificate or PCT International application having a filing date before that of the application on which priority is claimed:

PRIOR FOREIGN/PCT APPLICATION(S)

COUNTRY/OFFICE	APPLICATION NO.	DATE OF FILING	PRIORITY CLAIMED	
GB	9907216.7	March 29, 1999	☒YES	NO ☐
GB	9917565.5	July 27, 1999	☒YES	NO ☐
			☐YES	NO ☐
			☐YES	NO ☐

I hereby claim the benefit under 35 U.S.C. §119(e) of any United States provisional application(s) listed below.

PROVISIONAL APPLICATION NUMBER	DATE OF FILING

I hereby claim the benefit under 35 U.S.C. §120 of any United States application(s) or §365(c) of any PCT International application(s) designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of 35 U.S.C. §112, I acknowledge the duty to disclose material information as defined in 37 CFR §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

PRIOR U.S. APPLICATIONS OR PCT INTERNATIONAL APPLICATIONS DESIGNATING THE U.S. FOR BENEFIT UNDER 25 U.S.C. §120

		Status (check one)		
Application Serial No.	**Date of Filing**	**Patented**	**Pending**	**Abandoned**
		☐	☐	☐

(7) And I hereby appoint Arthur H. Seidel, Registration No. 15,979; Gregory J. Lavorgna, Registration No. 30,469; Daniel A. Monaco, Registration No. 30,480; Thomas J. Durling, Registration No. 31,349; John J. Marshall, Registration No. 29,671; Joseph R. Delmaster, Jr., Registration No. 38,399 and Robert E. Cannuscio, Registration No. 36,469, my attorneys or agents with full power of substitution and revocation, to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith.

Address all correspondence to Drinker Biddle & Reath LLP, One Logan Square, 18th & Cherry Streets, Philadelphia, PA 19103-6996. Address all telephone calls to **Daniel A. Monaco**, (215) 988-3312 (telefax: (215) 988-2757).

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

1-00

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FULL NAME OF SOLE OR SECOND INVENTOR

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Date: 29/10/01

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Residence: Portrush (City) Great Britain (State or Foreign Country) GBN

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